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VERIFICATION OF A TRANSLATION

I, Susan ANTHONY BA, ACIS,

Director of RWS Group Ltd, of Europa House, Marsham Way, Gerrards Cross, Buckinghamshire, England declare:

That the translator responsible for the attached translation is knowledgeable in the French language in which the below identified international application was filed, and that, to the best of RWS Group Ltd knowledge and belief, the English translation of the international application No. PCT/FR03/50003 is a true and complete translation of the above identified international application as filed.

I hereby declare that all the statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the patent application issued thereon.

Date: November 3, 2004

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For and on behalf of RWS Group Ltd

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POLYAMINO ACIDS FUNCTIONALIZED WITH  $\alpha$ -TOCOPHEROL,  
AND USES THEREOF, ESPECIALLY THERAPEUTIC USES

The present invention relates to novel materials based  
5 on biodegradable polyamino acids, which are useful  
especially for the vectorization of active principle(s)  
(AP).

The invention is also directed toward novel  
10 pharmaceutical, cosmetic, dietetic or plant-protection  
compositions based on these polyamino acids. These  
compositions may be of the type allowing the vectoriza-  
tion of AP and preferably being in the form of  
emulsions, micelles, particles, gels, implants or  
15 films.

The APs under consideration are advantageously biologi-  
cally active compounds that may be administered to an  
animal or human body via the oral, parenteral, nasal,  
20 vaginal, ocular, subcutaneous, intravenous, intramuscu-  
lar, intradermal, intraperitoneal, intracerebral,  
buccal, etc. route.

The APs more particularly concerned by the invention,  
25 but not limiting the invention thereto, are proteins,  
glycoproteins, peptides, polysaccharides, lipopoly-  
saccharides, oligonucleotides, polynucleotides and  
organic molecules. However, they may also be cosmetic  
products or plant-protection products, such as  
30 herbicides, insecticides, fungicides, etc.

In the field of vectorization of active principles,  
especially medicinal active principles, there is a  
need, in many cases:

- 35 • to protect them against degradation (hydrolysis,  
precipitation on site, enzymatic digestion, etc.)  
until they reach their site of action,  
• and/or to control their rate of release so as to

maintain a constant level over a given period,

- and/or to convey them (while protecting them) to the site of action.

5 To these ends, several types of polymers have been studied and some are even commercially available. Mention may be made, for example, of polymers of the polylactic, polylactic-glycolic, polyoxyethylene-oxypropylene, polyamino acid or polysaccharide type.  
10 These polymers constitute starting materials for manufacturing, for example, bulk implants, micro-particles, nanoparticles, vesicles, micelles or gels. Besides the fact that these polymers must be suitable for manufacturing such systems, they must also be  
15 biocompatible, nontoxic, nonimmunogenic and cost-effective, and they should be easy to eliminate from the body and/or biodegradable. As regards this last aspect, it is furthermore essential that the biodegradation in the body should generate nontoxic products.

20 By way of illustration of the prior art concerning polymers used as starting materials for making AP vectorization systems, various patents, patent applications or scientific articles are mentioned below.

25 Patent US 4 652 441 describes polylactide microcapsules encapsulating the hormone LH-RH. These microcapsules are produced by preparing a water-in-oil-in-water emulsion comprising an aqueous inner layer containing  
30 the hormone, a substance (gelatin) for fixing said hormone, a polylactide oily layer, and also an aqueous outer layer (polyvinyl alcohol). The AP is released over a period of more than 2 weeks after subcutaneous injection.

35 Patent US 6 153 193 describes compositions based on amphiphilic poly(oxyethylene)-poly(oxypropylene) micelles, for the vectorization of anticancer agents such

as adriamycin.

Akiyoshi et al. (J. Controlled Release 1998, 54, 313-320) describe pullulans that have been made hydrophobic by grafting cholesterol, and which form nanoparticles in water. These nanoparticles, which are capable of reversibly complexing with insulin, form stable colloidal suspensions.

10 Patent US 4 351 337 describes amphiphilic copolyamino acids based on leucine and glutamate, which may be used in the form of implants or microparticles for the controlled release of active principles. Said active principles are released over a very long period that  
15 depends on the rate of degradation of the polymer.

Patent US 4 888 398 describes polymers based on polyglutamate or polyaspartate, and optionally polyleucine, with pendent groups of alkyloxycarbonylmethyl type,  
20 placed randomly on the polyamino acid chain. These polyamino acids, grafted with side groups, e.g. methoxycarbonylmethyl, may be used in the form of biodegradable implants containing a sustained-release AP.

25 Patent US 5 904 936 describes nanoparticles obtained from a polyleucine-polyglutamate block polymer, which are capable of forming stable colloidal suspensions capable of spontaneously associating with biologically  
30 active proteins without denaturing them. Said proteins may then be released *in vivo* in a controlled manner, over a long period.

Patent application WO 00/30618 describes nanoparticles  
35 obtained from a poly(sodium glutamate)(polymethyl, ethyl, hexadecyl or dodecyl glutamate) block polymer, which are capable of forming stable colloidal suspensions capable of spontaneously associating with

biologically active proteins without denaturing them. Said proteins may then be released *in vivo* in a controlled manner, over a long period.

- 5 These amphiphilic copolyamino acids are modified by the presence of a hydrophobic alkyl side chain.

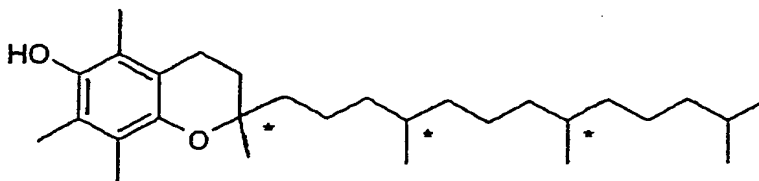
Patent US 5 449 513 describes amphiphilic block copolymers comprising a polyoxyethylene block and a  
10 polyamino acid block, for example poly( $\beta$ -benzyl-L-aspartate). These polyoxyethylene-polybenzylaspartate polymers form micelles that are capable of encapsulating hydrophobic active molecules such as adriamycin or indomethacin.

15

Patent application WO 99/61512 describes polylysines and polyornithines functionalized with a hydrophobic group (palmitic acid linked to polylysine or ornithine and a hydrophilic group (polyoxyethylene). These  
20 polymers, for example polylysine grafted with polyoxyethylene and palmitoyl chains, form, in the presence of cholesterol, vesicles capable of encapsulating doxorubicin or DNA.

- 25 It is moreover known practice to employ vitamin E derivatives, and more specifically  $\alpha$ -tocopherol, to construct AP vectorization systems.

Natural vitamin E consists of a mixture of compounds  
30 known as tocopherols (see Burton and Ingold, Acc. Chem. Res. 1986, 19, 194-201) and, in this mixture, the  $\alpha$ -tocopherol derivative is largely in majority amount. Vitamin E and some of its derivatives are nowadays used as a source of vitamin or as antioxidant in foods and  
35 cosmetic products. For these common uses, the vitamin E is found in its D- $\alpha$ -tocopherol form (its natural form) or in its D,L- $\alpha$ -tocopherol form (racemic and synthetic form). These two products are considered as being



As regards the vitamin E derivatives used in the field of vectorization of active principles, there is, at the present time and to the inventors' knowledge, no  $\alpha$ -tocopherol-based polymer product, with the exception of polymers of polyoxyethylene type, one end of which is grafted with  $\alpha$ -tocopherol-succinate groups. Polyethylene glycol grafted with  $\alpha$ -tocopherol succinate at the end of the chain (PEGylated vitamin E) is moreover available on the market, sold under the name TPGS 1000 by the company Eastman Chemical Ltd). This product, patented in 1954 (US 2 680 749), is nowadays used as a source of oral-route vitamin E. This polymer, and similarly  $\alpha$ -tocopherol-succinate and unmodified  $\alpha$ -tocopherol, have been proposed for the vectorization of active principles.

Patent US 5 869 703 describes similar compounds in which the polyoxyethylene chain comprises  $\alpha$ -tocopherol at one end and a (meth)acrylic residue at its other end. These  $\alpha$ -tocopherol derivatives are used to prepare stable amphiphilic vesicles (liposomes), used directly in cosmetic applications.

Patent application WO 00/71163 describes formulations based on polyethylene glycol grafted with  $\alpha$ -tocopherol-succinate at the end of the chain (TPGS 1000) and on  $\alpha$ -tocopherol, for the solubilization of paclitaxel  
5 (anticancer product). At the present time, the toxicity associated with the polyoxyethylene part is not known, and it is known that polyoxyethylene is not degraded *in vivo*. Furthermore, this compound contains only one  $\alpha$ -tocopherol unit per polymer chain and it has  
10 properties in solution similar to those of surfactants. In any case, the use of this product for vectorization would lead to sparingly stable polymer-active principle combinations.

15 Patent EP 0 243 446 describes the use of  $\alpha$ -tocopherol (hemi)succinate (organic acid derivative of  $\alpha$ -tocopherol) for the manufacture of vesicles, in combination with an amine salt. These vesicles may be used for the encapsulation of various active principles including  
20 small molecules, peptides and proteins. In general, it is indicated in said patent that the organic acid may be an amino acid or a polyamino acid. However, no details are given in this respect. Only  $\alpha$ -tocopherol (hemi)succinates are illustrated.

25 Thus, even though there are a great many technical solutions in the prior art, developed and proposed for the vectorization of medicinal active principles, meeting all the requirements is difficult to achieve,  
30 and remains unsatisfactory.

In this context, one of the essential objectives of the present invention is to provide a novel polymeric starting material, which may be used for AP  
35 vectorization and which can optimally satisfy all the specification details:

- O biocompatibility,
- O biodegradability,

O ability to be converted easily and economically into active principle vectorization particles,

O these particles themselves being capable:

- of forming stable aqueous colloidal suspensions,
- 5   • of readily associating with many active principles,
- and of releasing these active principles *in vivo*.

This objective, among others, is achieved by the  
10 present invention, which relates firstly to amphiphilic polyamino acids comprising aspartic units and/or glutamic units, characterized in that at least some of these units bear grafts comprising at least one  $\alpha$ -tocopherol unit.

15 These novel polymers have a biodegradable skeleton based on polyamino acids bearing side chains comprising  $\alpha$ -tocopherol. These polymers have association and/or encapsulation properties that are surprising in  
20 comparison with similar products and, what is more, they are readily degraded in the presence of enzymes.

The Applicant has, to its merit, had the idea of combining, in an entirely judicious and advantageous  
25 manner, particular biodegradable polyAsp and/or polyGlu polyamino acids with grafts based on  $\alpha$ -tocopherol (vitamin E), for the vectorization of AP.

For the purposes of the invention, the term "*polyamino*  
30 *acid*" covers not only oligoamino acids comprising from 2 to 20 amino acid units, but also polyamino acids comprising more than 20 amino acid units.

Preferably, the polyamino acids according to the  
35 present invention are oligomers or homopolymers comprising glutamic or aspartic amino acid repeating units or copolymers comprising a mixture of these two types of amino acid units, said units being partially

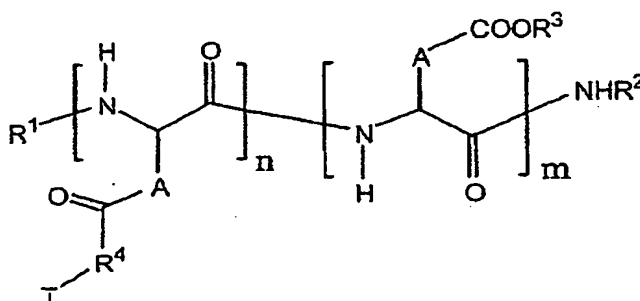


substituted with grafts comprising  $\alpha$ -tocopherol. The units under consideration in these polymers are amino acids having the D, L or D,L configuration and are linked via their  $\alpha$ - or  $\gamma$ -positions for the glutamate or glutamic unit and the  $\alpha$ - or  $\beta$ -position for the aspartic or aspartate unit.

The preferred amino acid units are those having the L configuration and a bond of  $\alpha$  type.

10

Even more preferably, the polyamino acids according to the invention correspond to the general formula (I) below:



(I)

15

in which:

- $R^1$  represents H, a C2 to C10 linear or C3 to C10 branched alkyl, benzyl or a terminal amino acid unit;
- 20 •  $R^2$  represents H, a linear C2 to C10 or branched C3 to C10 acyl group, or a pyroglutamate;
- $R^3$  is H or a cationic species preferably selected from the group comprising:
  - metallic cations advantageously chosen from the subgroup comprising sodium, potassium, calcium and magnesium,
  - 25 - organic cations advantageously chosen from the subgroup comprising:
    - amine-based cations,
    - 30 • oligoamine-based cations,
    - cations based on polyamine (polyethyleneimine

being particularly preferred),

- cations based on amino acid(s) advantageously chosen from the class comprising cations based on lysine or arginine,
- 5 - or cationic polyamino acids advantageously chosen from the subgroup comprising polylysine or oligo-lysine;
- $R^4$  represents a direct bond or a "spacer" based on 1 to 4 amino acid units;
- 10 • A independently represents a  $-CH_2-$  (aspartic unit) or  $-CH_2-CH_2-$  (glutamic unit) radical;
- $n/(n+m)$  is defined as the molar degree of grafting and ranges from 0.5 to 100 mol%;
- $n+m$  ranges from 3 to 1000 and preferably between 30
- 15 and 300;
- T represents an  $\alpha$ -tocopherol unit.

For these common uses, vitamin E is found in its D- $\alpha$ -tocopherol form (its natural form) or in its D,L- $\alpha$ -

20 tocopherol form (racemic and synthetic form). These two products are considered as being essentially nontoxic at doses considerably higher than therapeutic doses. In the context of the invention, these two forms of  $\alpha$ -tocopherol are preferred.

25

The  $\alpha$ -tocopherol is of natural or synthetic origin.

According to a first embodiment of the invention, the polyamino acids are  $\alpha$ -L-glutamate or  $\alpha$ -L-glutamic

30 homopolymers.

According to a second embodiment of the invention, the polyamino acids are  $\alpha$ -L-aspartate or  $\alpha$ -L-aspartic homopolymers.

35

According to a third embodiment of the invention, the polyamino acids are  $\alpha$ -L-aspartate/ $\alpha$ -L-glutamate or  $\alpha$ -L-aspartic/ $\alpha$ -L-glutamic copolymers.

Advantageously, the distribution of the aspartic and/or glutamic units bearing grafts comprising at least one  $\alpha$ -tocopherol unit is such that the polymers thus  
5 composed are either random, of block type or of multi-block type.

According to another mode of definition, the polyamino acids according to the invention have a molar mass of  
10 between 2000 and 100 000 g/mol and preferably between 5000 and 40 000 g/mol.

It is moreover preferable for the molar degree of grafting with  $\alpha$ -tocopherol of the polyamino acids  
15 according to the invention to be between 3% and 70% and preferably between 5% and 50%.

The polyamino acids of the invention are, remarkably, capable of being used in several ways depending on the  
20 degree of grafting. The methods for forming a polymer for the encapsulation of an active principle in the various forms targeted by the invention are known to those skilled in the art. For further details, reference may be made, for example, to these few  
25 particularly pertinent references:

*"Microspheres, Microcapsules and Liposomes; Vol. 1. Preparation and chemical applications"* Ed. R. Arshady, Citus Books 1999. ISBN: 0-9532187-1-6.

*"Sustained-Release Injectable Products"* Ed. J. Senior  
30 and M. Radomsky, Interpharm Press 2000. ISBN: 1-57491-101-5.

*"Colloidal Drug Delivery Systems"* Ed. J. Kreuter, Marcel Dekker, Inc. 1994. ISBN: 0-8247-9214-9.

*"Handbook of Pharmaceutical Controlled Release Technology"* Ed. D.L. Wise, Marcel Dekker, Inc. 2000. ISBN:  
35 0-8247-0369-3.

The polyamino acids are also extremely advantageous as

a result of the fact that, at a relatively low degree of grafting of about from 3% to 10%, they form in water at pH 7.4 (for example with a phosphate buffer) colloidal suspensions or gels depending on the polymer concentration. Furthermore, the polyamino acid particles forming the dispersed phase of the colloidal suspension can readily associate with active principles such as proteins, peptides or small molecules. The preferred forming operation is that described in patent application WO 00/30618 by the Applicant, which consists in dispersing the polymer in water and incubating the solution in the presence of an AP. This solution can then be filtered through a 0.2  $\mu$ m filter and then injected directly into a patient.

Above a degree of grafting of 10%, the polymer can form microparticles capable of associating or of encapsulating APs. In this context, the forming of the microparticles may take place by codissolving the AP and the polymer in a suitable organic solvent, and the mixture is then precipitated from water. The particles are then recovered by filtration and can then be used for an oral administration (in the form of a gel capsule, in compacted and/or coated form, or alternatively in a form dispersed in an oil) or parenterally after redispersing in water.

At degrees of grafting of greater than 30%, the redispersion of the polymer in aqueous phase becomes more difficult due to the smaller amount of ionizable carboxylate functions, and the polymer precipitates. In this case, the polymer can be dissolved in a biocompatible solvent such as N-methylpyrrolidone or a suitable oil such as Miglyol<sup>®</sup>, and then injected intramuscularly or subcutaneously or into a tumor. The diffusion of the solvent or of the oil results in precipitation of the polymer at the site of injection and thus forms a deposit. These deposits then provide controlled release

by diffusion and/or erosion and/or hydrolytic or enzymatic degradation of the polymer.

5 In general, the polymers of the invention, in neutral or ionized form, may be used alone or in a liquid, solid or gel composition and in an aqueous or organic medium.

10 It should be understood that the polymer based on polyamino acids contains carboxylic functions that are either neutral (COOH form) or ionized, depending on the pH and composition. For this reason, the solubility in an aqueous phase depends directly on the content of free COOH (not grafted with vitamin E) and on the pH.

15 In aqueous solution, the counter-cation may be a metallic cation such as sodium, calcium or magnesium, or an organic cation such as triethanolamine, tris(hydroxymethyl)aminomethane or a polyamine such as polyethyleneimine.

20

The polymers of the invention are obtained via methods known to those skilled in the art. The polyamino acids may be obtained in at least two ways:

- grafting of  $\alpha$ -tocopherol onto a polyamino acid, or
- 25 • polymerization of NCA derivatives of  $\alpha$ -tocopherol, followed by a selective hydrolysis.

30 In the first case, a polyamino acid, homopolyglutamate, homopolyaspartate or a glutamate/aspartate copolymer, in block, multiblock or random form, is prepared, for example, according to standard methods.

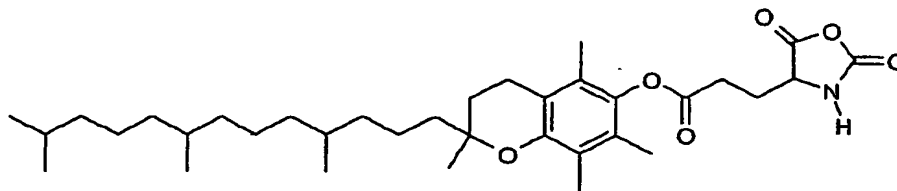
To obtain polyamino acids of  $\alpha$  type, the technique most commonly used is based on the polymerization of

35 N-carboxyamino acid anhydrides (NCA), described, for example, in the article "Biopolymers, 1976, 15, 1869" and in the book by H.R. Kricheldorf "alpha-Aminoacid-N-carboxy Anhydride and related Heterocycles", Springer

Verlag (1987). The NCA derivatives are preferably NCA-O-Me, NCA-O-Et or NCA-O-Bz derivatives (Me = methyl, Et = ethyl and Bz = benzyl). The polymers are then hydrolyzed under suitable conditions to give the polymer in its acid form. These methods are inspired from the description given in patent FR 2 801 226 from the Applicant. A certain number of polymers that may be used according to the invention, for example, of poly( $\alpha$ -L-aspartic), poly( $\alpha$ -L-glutamic), poly( $\alpha$ -D-glutamic) and poly( $\gamma$ -L-glutamic) type of variable masses are commercially available. The polyaspartic of  $\alpha$ - $\beta$  type is obtained by condensing aspartic acid (to obtain a poly-succinimide), followed by a basic hydrolysis (see Tomida et al. Polymer 1997, 38, 4733-36).

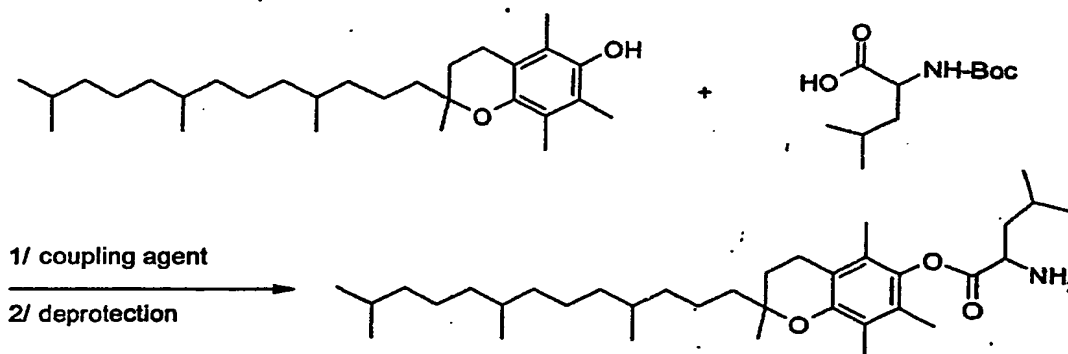
15 The coupling of  $\alpha$ -tocopherol with an acid function is readily performed by reacting the polyamino acid with vitamin E in the presence of a carbodiimide as coupling agent and, preferably, a catalyst such as 4-dimethyl-  
20 aminopyridine, and in a suitable solvent such as dimethylformamide (DMF), N-methylpyrrolidone (NMP) or dimethyl sulfoxide (DMSO). The carbodiimide is, for example, dicyclohexylcarbodiimide or diisopropylcarbo-  
diimide. The degree of grafting is controlled  
25 chemically via the stoichiometry of the constituents and reagents, or the reaction time.

In the second case, an NCA derivative of  $\alpha$ -tocopherol having the structure below is synthesized. The synthesis is analogous to that described for stearyl glutamate N-carboxyanhydride by Poché et al. *Macromolecules* 1995, 28, 6745-53.



The NCA derivative of  $\alpha$ -tocopherol-glutamate is then copolymerized with, for example, benzylglutamate NCA and, to obtain the glutamate or glutamic functions, a selective hydrolysis reaction of the benzyl functions is performed in a mixture of trifluoroacetic acid and hydrobromic acid at room temperature. It should be noted that this second synthetic route makes it possible easily to prepare random, block or multiblock copolymers simply by modifying the order of addition of the monomers.

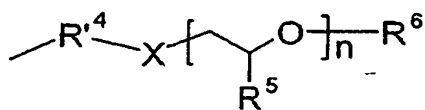
The coupling of vitamin E via a spacer consisting of 1 to 4 amino acids may be performed via successive reactions of vitamin E with suitably protected amino acids, which are then deprotected to have a graftable amine function on the polymer, or by reaction with an oligopeptide. For example, the synthesis of an  $\alpha$ -tocopherol with a leucine unit is performed according to a general method well known to those skilled in the art and according to the following scheme:



It should be noted that the direct grafting of  $\alpha$ -tocopherol onto the polymer takes place via an ester function, whereas in the case of the presence of a spacer based on amino acid(s), it takes place via an amide function. As for the preparation of an ester bond, the amide bond may be formed in the same manner using a standard coupling agent such as a dialkylcarbo-

diimide.

According to one variant of the invention, the poly-  
amino acids with which it is concerned not only bear  
5  $\alpha$ -tocopherol grafts but also, per molecule, at least  
one graft of polyalkylene glycol type linked to a  
glutamate and/or aspartate unit, and preferably of  
formula (II) below:



(II)

10

in which

$\text{R}^4$  represents a direct bond or a "spacer" based on  
1 to 4 amino acid units;

15 X is a hetero atom chosen from the group comprising  
oxygen, nitrogen and sulfur;

$\text{R}^5$  and  $\text{R}^6$  independently represent H or a linear C1 to  
C4 alkyl.

n ranges from 3 to 1000.

20

Preferably, the polyalkylene glycol is a polyethylene  
glycol.

According to another preferred characteristic of the  
25 invention, the molar percentage of grafting of the  
polyalkylene glycol ranges from 1% to 30%.

The grafting of these pendent side groups (II) is  
performed in a manner that is known per se and  
30 according to techniques within the capability of a  
person skilled in the art, for example by forming  
amide, ester or thioester bonds with the carboxyls of  
glutamate and/or aspartate monomers. These techniques  
may especially be those used for the grafting of  
35  $\alpha$ -tocopherol onto a polyamino acid skeleton, said



techniques being described in the present patent application.

5 According to another of its aspects, the invention is directed toward a pharmaceutical, cosmetic, dietetic or plant-protection composition comprising at least one of the polyamino acids as defined above.

10 According to one advantageous arrangement of the invention, this composition comprises, besides  $\alpha$ -tocopherol, at least one active principle, which may be a therapeutic, cosmetic, dietetic or plant-protection active principle.

15 Preferably, the active principle is a protein, a glycoprotein, a polysaccharide, a liposaccharide, an oligonucleotide, a polynucleotide or a peptide.

20 Even more preferably, the active principle is a hydrophobic, hydrophilic or amphiphilic organic "small molecule".

According to the present description, the term "small molecule" especially denotes nonprotein molecules.

25 This composition may be in the form of nanoparticles, microparticles, solutions, emulsions, suspensions, gels, micelles, implants, powders or films.

30 According to one of its particularly preferred forms, the composition, containing or not containing active principle(s), is a stable colloidal suspension of nanoparticles and/or microparticles and/or micelles of polyamino acids, in an aqueous phase.

35 The composition according to the invention, when it is pharmaceutical, may be administered via the oral, parenteral, nasal, vaginal, ocular, subcutaneous,

intravenous, intramuscular, intradermal, intraperitoneal, intracerebral or buccal route.

It may also be envisioned for the composition to be in the form of a solution in a biocompatible solvent, capable of being injected subcutaneously, intramuscularly or into a tumor.

According to another variant, the composition according to the invention is formulated such that it is injectable and such that it is capable of forming a deposit at the site of injection.

The invention is also directed toward compositions comprising polyamino acids according to the invention and active principles, and which are capable of being used for the preparation:

- of medicinal products, in particular for oral, nasal, vaginal, ocular, subcutaneous, intravenous, intramuscular, intradermal, intraperitoneal or intracerebral administration, the active principles of these medicinal products possibly being, especially, proteins, glycoproteins, proteins linked to one or more polyalkylene glycol chains {for example polyethylene glycol (PEG), in which case they are referred to as "PEGylated" proteins}, peptides, polysaccharides, liposaccharides, oligonucleotides, polynucleotides and hydrophobic, hydrophilic or amphiphilic organic small molecules;
- and/or nutrients;
- and/or cosmetic or plant-protection products.

According to yet another of its aspects, the invention is directed toward a process for preparing:

- medicinal products, in particular for oral, nasal, vaginal, ocular, subcutaneous, intravenous, intramuscular, intradermal, intraperitoneal or intracerebral administration, the active principles of

these medicinal products possibly being, especially, proteins, glycoproteins, proteins linked to one or more polyalkylene glycol chains (for example polyethylene glycol (PEG), in which case they are referred to as "PEGylated" proteins), peptides, polysaccharides, liposaccharides, oligonucleotides, polynucleotides and hydrophobic, hydrophilic or amphiphilic organic small molecules;

- and/or nutrients;

- and/or cosmetic or plant-protection products;

this process being characterized in that it consists essentially in using at least one polyamino acid as defined above and/or the composition itself also described above.

As indicated above, the techniques for associating one or more APs with the  $\alpha$ -tocopherol-grafted polyamino acids according to the invention are described especially in patent application WO 00/30618.

The invention also relates to a therapeutic treatment method that consists essentially in administering the composition as described in the present description, via the oral, parenteral, nasal, vaginal, ocular, subcutaneous, intravenous, intramuscular, intradermal, intraperitoneal, intracerebral or buccal route.

According to one particular embodiment, the therapeutic treatment method consists essentially in using a composition as described above in the form of a solution in a biocompatible solvent, and then injecting it subcutaneously, intramuscularly or into a tumor, preferably such that it forms a deposit at the site of injection.

As examples of APs that may be associated with the polyamino acids according to the invention, whether or not they are in the form of (nano or micro)particles,

mention may be made of:

- proteins such as insulin, interferons, growth hormones, interleukins, erythropoietin or cytokines;
- peptides such as leuprolide or cyclosporin;
- 5 ○ small molecules such as those belonging to the anthracyclin, taxoid or camptothecin family;
- and mixtures thereof.

The invention will be understood more clearly and its  
10 advantages and implementation variants will emerge more clearly from the examples that follow, which describe the synthesis of  $\alpha$ -tocopherol-grafted polyamino acids, their conversion into an AP vectorization system (stable aqueous suspension of nanoparticles) and  
15 demonstration of the capacity of such a system to associate with APs (small organic molecules, proteins, etc.) to form pharmaceutical compositions.

**Example 1: Polymer P1**

20 ***Synthesis of a polyglutamate grafted with  $\alpha$ -tocopherol of synthetic origin***

The  $\alpha$ -L-polyglutamate polymer, with a mass equivalent to about 10 000 relative to a polyoxyethylene standard,  
25 is obtained by polymerization of NCAGluOMe, followed by a hydrolysis, as described in patent application FR 2 801 226. 5.5 g of this  $\alpha$ -L-polyglutamate polymer are dissolved in 92 ml of dimethylformamide (DMF), by heating at 40°C for 2 hours. Once the polymer has  
30 dissolved, the temperature is allowed to return to 25°C and 1.49 g of D,L, $\alpha$ -tocopherol (> 98%, obtained from Fluka®) predissolved in 6 ml of DMF, 0.09 g of 4-dimethylaminopyridine predissolved in 6 ml of DMF and 0.57 g of diisopropylcarbodiimide predissolved in 6 ml  
35 of DMF are successively added. After stirring for 8 hours at 25°C, the reaction medium is poured into 800 ml of water containing 15% sodium chloride and hydrochloric acid (pH 2). The precipitated polymer is

then recovered by filtration, washed with 0.1N hydrochloric acid and then washed with water. The polymer is then redissolved in 75 ml of DMF and then reprecipitated from water containing, as previously,  
5 salt and acid at pH 2. After washing twice with water, the polymer is washed several times with diisopropyl ether. The polymer is then oven-dried under vacuum at 40°C. A yield of about 85% is obtained.

10 The degree of grafting estimated by proton NMR is about 7.8% and an HPLC analysis reveals a residual tocopherol content of less than 0.3%.

Mw (measured by GPC, eluting with NMP) = 17 500 g/mol (as polymethyl methacrylate equivalent).

15

**Examples 2, 3, 4 and 5: Synthesis of polymers P2, P3, P4 and P5**

Polymers containing variable amounts of tocopherol are prepared in the same manner.

20

Table 1:

Polymer	$\alpha$ -Tocopherol	Degree of grafting
P2	Synthetic: D,L	5.2%
P3	Synthetic: D,L	12.8%
P4	Synthetic D,L	20.0%
P5	Synthetic D,L	50.0%

In all cases, the amount of tocopherol effectively  
25 grafted was confirmed by NMR.

**Example 6: Polymer P6**

***Synthesis of a polyglutamate grafted with  $\alpha$ -tocopherol of natural origin***

30

In a similar manner, the polymer P6 is synthesized with 7.3% of D- $\alpha$ -tocopherol of natural origin (at a purity of 98.5%, obtained from the company ADM France).

The molar mass is 17 400 (GPC NMP, PMMA eq).

**Example 7: Analysis of the polymers in aqueous solution**

The polymers are dissolved in a phosphate-buffered  
5 saline at pH 7.4 at concentrations ranging from 10 to  
40 mg/ml, and the pH is adjusted to 7.4 by adding 0.1N  
sodium hydroxide. The dissolution is observed visually.

Table 2: Solubility in saline water at pH 7.4

10

Polymer	Degree of grafting	Concentration	Appearance
P1 (D,L)	7.8%	10 to 30 mg/ml	soluble and clear
P3 (D,L)	12%	10 mg/ml	very fine precipitate
P4 (D,L)	20%	10 mg/ml	very fine precipitate
P6 (D)	7.3%	10 to 30 mg/ml	soluble and clear

Phosphate buffer: 0.01M phosphate, 0.0027M KCl and  
0.137M NaCl.

An observation by electron transmission of the clear  
15 solutions of the polymer P1 deposited on a support  
shows the existence of nanoparticles of 15 to 25 nm. A  
comparative analysis of the solutions of polymer P1, P6  
and  $\alpha$ -tocopherol succinate at 15 mg/ml in water at pH  
7.4 (phosphate buffer) reveals that only the  $\alpha$ -toco-  
20 pherol succinate develops a milky solution characteris-  
tic of vesicles, as described in patent EP 0 243 446.

**Example 8: Adsorption of a dye onto the polymer P1**

According to one of the subjects of the invention, the  
25 polymers may be used in the form of a colloidal  
suspension in water and associated with an active  
principle. For this application, it is demonstrated in  
the experiment below that with certain polymers,  
especially those with a degree of grafting of about  
30 from 5% to 10% tocopherol, the adsorption capacity is  
greater than that of a similar compound of the prior  
art.

For this study, the polymer P1 was compared with a similar polymer containing a dodecanol chain grafted onto a polyglutamate. This polymer is described in  
5 patent WO 00/30618.

The study is performed in the following manner: the polymers are dissolved in an aqueous solution at pH 7 (phosphate buffer) and 5 mg of the dye known as Orange  
10 OT (Rn CAS: 2646-17-5) are added. The solutions are left in an ultrasonic bath for 1 hour to achieve the association. The solutions are then centrifuged to remove the nonassociated dye, and the optical density is measured at the  $\lambda_{\text{max}}$  of the dye, which is at 495 nm.

15

Table 3:

Polymer	Degree of grafting	Polymer concentration	Normalized OD
P1 ( $\alpha$ -tocopherol)	7.8 mol%	13.8 mg/ml	1
Comparative polymer* (dodecanol)	15 mol%	17.3 mg/ml	0.45

\*WO 00/30618

20 It is found that at a molar degree of grafting of less than a half and at a slightly smaller mass concentration of polymer, the polymer P1 has a much higher capacity for association of the dye Orange OT.

25 **Example 9: Synthesis of the polymer P7**

***Synthesis of a polyglutamate containing an  $\alpha$ -tocopherol leucine graft***

The  $\alpha$ -tocopherol leucine derivative is first  
30 synthesized, in the following manner.

D,L- $\alpha$ -Tocopherol (4.3 g) is reacted with BOC-leucine (2.3 g) in 15 ml of dichloromethane in the presence of 4-dimethylaminopyridine (244 mg) and diisopropylcarbodiimide (1.5 g). After 2 hours at 30°C the product is purified by filtration on a column of silica. 5 g of the product  $\alpha$ -tocopherol leucine BOC are obtained (77% yield). Its structure is confirmed by NMR spectroscopy. The deprotection of the product is performed in trifluoroacetic acid at a temperature of between 5 and 10°C for 1 hour. After purification by filtration through silica, 3.3 g of the desired product are isolated (78% yield). Its structure is confirmed by NMR spectroscopy.

The grafting reaction on a polyglutamic acid is then performed under the same conditions as in example 1, with a degree of grafting of 7%. The structure of the polymer and the degree of grafting were confirmed by NMR spectroscopy.

20

#### **Example 10: Synthesis of polymer P8**

Synthesis of a polyglutamate containing an  $\alpha$ -tocopherol graft and a polyoxyethylene glycol graft.

25 A grafting reaction is performed as in example 1, with 11 mol% of  $\alpha$ -tocopherol and 2 mol% of an amino methoxypolyethylene glycol of formula  $\text{MeO}(\text{CH}_2\text{CH}_2\text{O})_a\text{CH}_2\text{CH}_2\text{NH}_2$  and of molar mass 3000 (product obtained from the company Shearwaters). The polymer in its acid form is obtained in a yield of 72%. Proton NMR confirms a degree of grafting with  $\alpha$ -tocopherol of 10.9% and with polyethylene glycol of 1.9%.

30

#### **Example 11: Adsorption of insulin**

35 A solution containing 1 mg of polymer P1 and 7 mg of insulin at pH 7.0 in 1 ml of water is prepared and is left to incubate for 2 hours. The suspension is then ultrafiltered (10 000xG, 20 minutes with a 100 KDa



threshold). The free insulin in the filtrate is assayed by HPLC and the amount of associated insulin is deduced by difference. A degree of association of greater than 95% relative to the insulin employed is measured. Under  
5 the same conditions, the comparative polymer of example 8 allows 40% association. The adsorption capacity of polymer P1 is thus greater.

**Example 12: *in-vitro* degradation of polymer P1 in the  
10 presence of enzymes**

Polymer P1 is dissolved at pH 7.5 (phosphate buffer and 10 mM of calcium cation) and at a concentration of 20 mg/ml. 0.1 ml of protease (solution of 10 mg/ml) is added and the degradation is monitored by aqueous GPC.

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A relatively rapid degradation is found, with a half-life time of the initial polymer of about 100 minutes.